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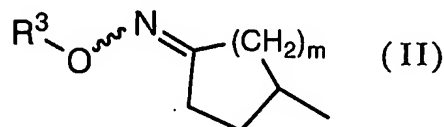
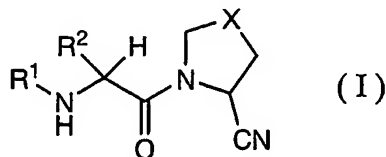
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(54) Title: DPP-IV INHIBITOR



(57) Abstract: A compound of the formula (I) or a pharmaceutically acceptable salt thereof: [wherein X is CH₂#191, O or S, R₁ is the moiety represented by the formula (II), etc.: [wherein R₃ is lower alkyl, etc., m is integer of 1 to 3], R₂ is hydrogen, etc.]. Compound of formula (I) inhibits DPP-IV activity. They are therefore useful in the treatment of conditions mediated by DPP-IV, such as NIDDM.

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DESCRIPTION

DPP-IV INHIBITOR

TECHNICAL FIELD

5 This invention relates to the compounds and pharmaceutically acceptable salts thereof which inhibit dipeptidyl peptidase-IV (DPP-IV).

BACKGROUND ART

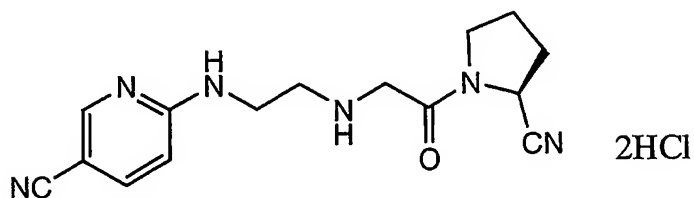
10 It is known that DPP-IV has various physiological functions in living body, especially has the action which inactivates Glucagon-like peptide-1 (GLP-1) by cleaving the terminal dipeptide (His-Ala). And the resultant peptide is the receptor antagonist of GLP-1 and totally
15 reduces the activity of GLP-1.

 This GLP-1 has very important role in sugar metabolism. For example, (1) GLP-1 intensifies the secretion of insulin, (2) express genes which are indispensable for the secretion of insulin, (3) stimulate proliferation of
20 β -cell, (4) suppresses secretion of glucagon, (5) suppresses the function about secretion and motility of digestive organs (especially, peristalsis), and (6) suppresses appetite. That is, GLP-1 restricts food ingestion, postpones the process of digestion and
25 absorption, and raised the use of the sugar in blood.

 Therefore, the inhibitor of DPP-IV can maintain the activity of GLP-1, so it is expected as a medicine to treat and prevent various diseases, especially non-insulin dependent diabetes mellitus (NIDDM).

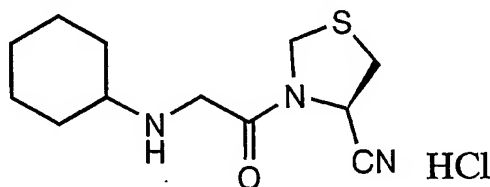
30 Hitherto, such inhibitors of DPP-IV are known so far.

For example, in US 6,011,155, 2-cyanopyrrolidine compounds like following are disclosed.



Pyrrolidine, 1-[[2-[(5-cyano-2-pyridinyl)amino]ethyl]amino]acetyl-2-cyano, (S)-, dihydrochloride

In US 6,110,949, 4-cyanothiazolidine compounds like
5 following are disclosed.



3-[(Cyclohexyl)amino]acetyl-4-cyano-(R)-thiazolidine monohydrochloride

DISCLOSURE OF INVENTION

Under the above situation, the inventors of this
10 invention found that the introduction of the oxime
derivative group at appropriate position of the compound
results in remarkable improvement of the activity to
inhibit DPP-IV. And the inventors completed this
invention.

15 Accordingly, this invention relates to DPP-IV
inhibitor. More particularly, this invention relates to
DPP-IV inhibitor useful for treating and/or preventing
the medical conditions mediated by DPP-IV, more
particularly useful for treating and/or preventing

altered glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes mellitus (IDDM and NIDDM), diabetic neuropathy, nephropathy, and secondary diseases in mammals caused by diabetes mellitus.

5 That is, one object of this invention is to provide novel compounds and pharmaceutically acceptable salts thereof, of which activity to inhibit DPP-IV is remarkably improved against known compounds.

10 Another object of this invention is to provide a medicament and pharmaceutical composition containing the compound and/or pharmaceutically acceptable salts thereof as an active ingredient.

15 A further object of this invention is to provide an inhibitor of DPP-IV consisting of the compound and/or pharmaceutically acceptable salts thereof.

20 A further object of this invention is to provide a method for treating and/or preventing the medical conditions comprising to administer an effective amount of the compounds and/or pharmaceutically acceptable salts thereof.

 A further object of this invention is to provide a use of the compounds and pharmaceutically acceptable salts thereof as medicaments.

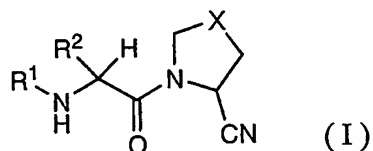
25 A further object of this invention is to provide the compounds and pharmaceutically acceptable salts thereof which are useful for the manufacture of medicaments for treating and/or preventing the medical conditions mediated by DPP-IV

30 A further object of this invention is to provide the commercial package comprising the pharmaceutical

composition containing the compound and a written matter associated therewith, wherein the written matter states that the compound can or should be used for preventing and/or treating the medical conditions mediated by DPP-IV.

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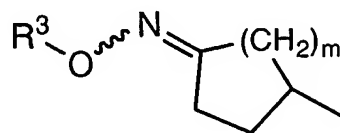
The compounds of this invention can be represented by the following formula (I):



[wherein

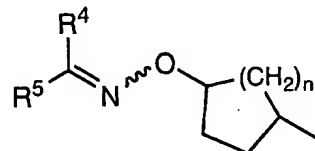
10 X is CH₂, O or S,

R¹ is hydrogen, lower alkyl, the moiety represented



by the formula:

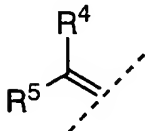
[wherein R³ is lower alkyl, aryl (which may have 1 to 3 substituent(s) selected from the group described later), heteroaryl (which may have 1 to 3 substituent(s) selected from the group described later), aryl-(lower alkyl) (which may have 1 to 3 substituent(s) selected from the group described later on the aryl group), or heteroaryl-(lower alkyl) (which may have 1 to 3 substituent(s) selected from the group described later on the heteroaryl group), m is integer of 1 to 3], or



the moiety represented by the formula:

[wherein R⁴ is hydrogen or lower alkyl, R⁵ is lower alkyl, aryl (which may have 1 to 3 substituent(s) selected

from the group described later), or heteroaryl (which may have 1 to 3 substituent(s) selected from the group described later), or the partial structure:



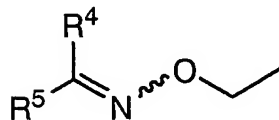
may form cycloalkyl, n is integer of 1 to

5 3],

R² is hydrogen, the moiety represented by the formula:



[wherein R³ represents the same definition defined above], or the moiety represented by the formula:



10

[wherein R⁴ and R⁵ represent the same definition defined above],

the substituent(s) is(are) selected from the group consisting of lower alkyl, lower alkoxy, halogen, cyano, nitro, amino and hydroxy,

15

provided that both of R¹ and R² are not hydrogen at the same time; and either R¹ and R² is hydrogen, or when R¹ is lower alkyl, R² is not hydrogen.]

20

In the above and subsequent description of the present specification, suitable examples of the various definitions to be included within the scope of the invention are explained in detail in the following.

25

The term "lower" is intended to mean a group having

1 to 6 carbon atom(s), unless otherwise provided.

So, the "lower alkyl" means a straight or branched chain aliphatic hydrocarbon, such as methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tert-butyl, pentyl, hexyl, and the like, and it is preferably C1-C4 alkyl, more preferably C1-C2 alkyl, most preferably methyl.

The "aryl" means C6-C10 aromatic hydrocarbon group, such as phenyl, indenyl, naphthyl, and the like, and it is preferably phenyl. And the "aryl" may have 1 to 3 substituent(s), the number of substituent is preferably 1 or 2, more preferably 1, in case that "aryl" has plural substituents, they may be the same or different each other, but, needless to say, "aryl" may not have substituent.

The "heteroaryl" means 5- or 6-membered aromatic heterocyclic group which contains at least one hetero atom such as nitrogen, oxygen, sulfur atom, preferably contains nitrogen atom. The "heteroaryl" may include 5-membered heteroaryl group such as pyrrolyl, imidazolyl, pyrazolyl, thienyl, furyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl; and 6-membered heteroaryl group such as pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, or the like, and is preferably 6-membered heteroaryl, more preferably pyridyl. And the "heteroaryl" may have 1 to 3 substituent(s), the number of substituent is preferably 1 or 2, more preferably 1, in case that "heteroaryl" has plural substituents, they may be the same or different each other, but, needless to say, "heteroaryl" may not have substituent.

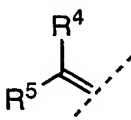
The "aryl-(lower alkyl)" means the "lower alkyl" group substituted by "aryl" and include benzyl,

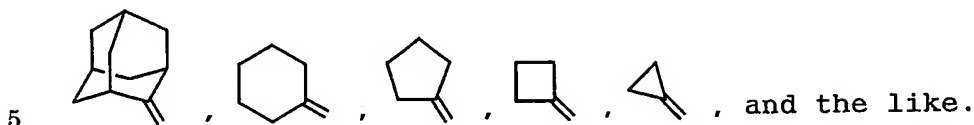
naphthylmethyl, 1-phenethyl, 2-phenethyl,
1-naphthylmethyl, 2-naphthylmethyl, 3-phenylpropyl,
3-naphthylpropyl, 4-phenylbutyl, 4-naphthylbutyl,
6-phenylhexyl, 6-naphthylhexyl and the like, and it is
5 preferably phenyl-(lower alkyl), more preferably
phenyl-(C1-C4 alkyl), more preferably phenyl-(C1-C2
alkyl), most preferably benzyl. And the "aryl-(lower
alkyl)" may have 1 to 3 substituent(s) on the aryl group,
the number of substituent is preferably 1 or 2, more
10 preferably 1, in case that "aryl-(lower alkyl)" has plural
substituents, they may be the same or different each other,
but, needless to say, "aryl-(lower alkyl)" may not have
substituent.

The "heteroaryl-(lower alkyl)" means the "lower
15 alkyl" group substituted by "heteroaryl" such as
pyrrolylmethyl, thienylmethyl, furylmethyl,
pyridylmethyl, oxazolylmethyl, thiazolylmethyl,
pyrrolylethyl, pyridylethyl, oxazolylethyl,
thiazolylethyl, pyridylbutyl, oxazolylbutyl,
20 thiazolylbutyl or the like, and it is preferably
(6-membered heteroaryl)-(C1-C4 alkyl), more preferably
(6-membered heteroaryl)-(C1-C2 alkyl), most preferably
pyridylmethyl. And the "heteroaryl-(lower alkyl)" may
have 1 to 3 substituent(s) on the heteroaryl group, the
25 number of substituent is preferably 1 or 2, more preferably
1, in case that "heteroaryl-(lower alkyl)" has plural
substituents, they may be the same or different each other,
but, needless to say, "heteroaryl-(lower alkyl)" may not
have substituent.

30 The "cycloalkyl" means C3-C10 cycloalkyl group, such

as cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, norbornyl, adamantyl, and the like, and it is preferably cyclohexyl or adamantyl. So, the partial

structure:  is exemplified



The "lower alkoxy" means a straight or branched chain aliphatic hydrocarbon oxy group, such as methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, tert-butoxy, pentoxy, hexoxy, and the like, and it is preferably C1-C4
10 alkoxy, more preferably C1-C2 alkoxy, most preferably methoxy.

The "halogen" may include a fluorine atom, a chlorine atom, a bromine or a iodine atom, more preferably a chlorine atom.

15

The compounds of formula (I) may contain one or more asymmetric centers and thus they can exist as enantiomers or diastereoisomers. This invention includes both mixtures and separate individual isomers. However, in
20 the portion of 2-cyanopyrrolidine, (2S) isomer is more preferable, and in the portion of 2-cyanooxazolidine and 2-cyanothiazolidine, (4R) isomer is more preferable.

Oxime derivatives may have two kind of geometrical isomers, that is, (syn) form and (anti) form, this
25 invention includes both isomers.

The compounds of the formula (I) may also exist in

tautomeric forms and this invention includes both mixtures and separate individual tautomers.

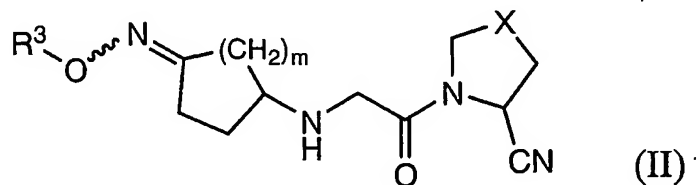
The compounds of the formula (I) and its salts may be in a form of a solvate such as hydrate, which is included
5 within the scope of the present invention.

Also included in the scope of invention are radiolabelled derivatives of compounds of formula (I) which are suitable for biological studies.

The compounds of this invention may be converted to
10 salt according to a conventional method. Suitable salts of the compounds (I) are pharmaceutically acceptable conventional non-toxic salts such as an organic acid salt (e.g., acetate, maleate, tartrate, methanesulfonate, benzenesulfonate, formate, toluenesulfonate,
15 trifluoroacetate, etc.), an inorganic acid salt (e.g., hydrochloride, hydrobromide, sulfate, phosphate, etc.), a salt with an amino acid (e.g., arginine, aspartic acid, glutamic acid, etc.), or the like, and the preferable salt is hydrochloride or trifluoroacetate salt.

20

The Compound (I) may preferably include;
a compound of the formula (II)



[wherein

25 X is CH_2 , O or S,

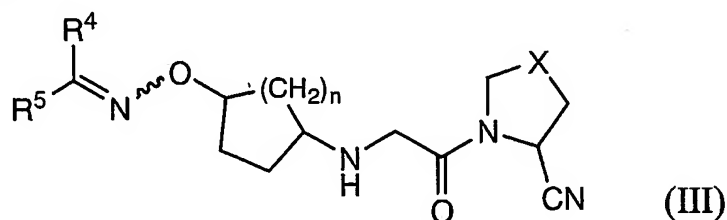
R^3 is lower alkyl, aryl (which may have 1 to 3 substituent(s) selected from the group described later),

heteroaryl (which may have 1 to 3 substituent(s) selected from the group described later), aryl-(lower alkyl) (which may have 1 to 3 substituent(s) selected from the group described later on the aryl group), or heteroaryl-(lower alkyl) (which may have 1 to 3 substituent(s) selected from the group described later on the heteroaryl group),

m is integer of 1 to 3

the substituent(s) is selected from the group consisting of lower alkyl, halogen, cyano, nitro, amino and hydroxyl,

a compound of the formula (III)



[wherein

X is CH₂, O or S,

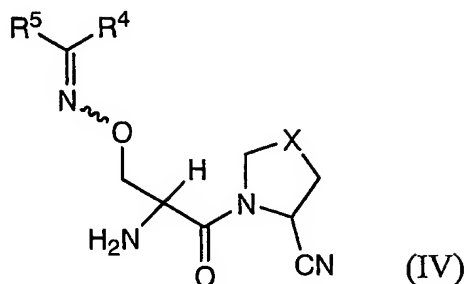
R⁴ is hydrogen or lower alkyl,

R⁵ is lower alkyl, aryl (which may have 1 to 3 substituent(s) selected from the group described later), or heteroaryl (which may have 1 to 3 substituent(s) selected from the group described later),

n is integer of 1 to 3

the substituent(s) is(are) selected from the group consisting of lower alkyl, halogen, cyano, nitro, amino and hydroxyl,

a compound of the formula (IV)



[wherein

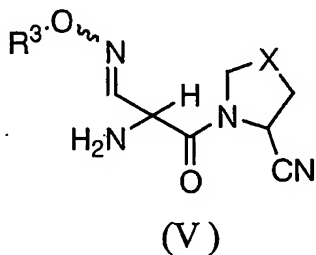
X is CH₂, O or S,

5 R⁴ is hydrogen or lower alkyl,

R⁵ is lower alkyl, aryl (which may have 1 to 3 substituent(s) selected from the group described later), or heteroaryl (which may have 1 to 3 substituent(s) selected from the group described later),

10 the substituent(s) is(are) selected from the group consisting of lower alkyl, halogen, cyano, nitro, amino and hydroxy] , and

a compound of the formula (V)



15 [wherein

X is CH₂, O or S,

R³ is lower alkyl, aryl (which may have 1 to 3 substituent(s) selected from the group described later), heteroaryl (which may have 1 to 3 substituent(s) selected from the group described later), aryl-(lower alkyl) (which may have 1 to 3 substituent(s) selected from the group

20

described later on the aryl group), or heteroaryl-(lower alkyl) (which may have 1 to 3 substituent(s) selected from the group described later on the heteroaryl group),

the substituent(s) is(are) selected from the group
5 consisting of lower alkyl, halogen, cyano, nitro, amino and hydroxyl.

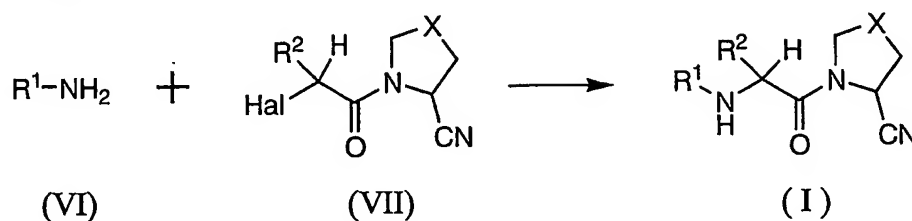
And in the each definition of the compound formula(I), preferably,

- 10 (1) X is CH₂ or S,
(2) X is CH₂,
(3) R³ is lower alkyl, aryl-(lower alkyl) (which may have 1 to 3 substituent(s) on the aryl group), or heteroaryl-(lower alkyl) (which may have 1 to 3
15 substituent(s) on the heteroaryl group),
(4) R³ is C1-C4 alkyl, aryl-methyl (which may have 1 to 3 substituent(s) on the aryl group), or heteroaryl-methyl (which may have 1 to 3 substituent(s) on the heteroaryl group),
20 (5) R³ is C1-C4 alkyl, benzyl (which may have 1 to 3 substituent(s) on the phenyl group), or pyridylmethyl (which may have 1 to 3 substituent(s) on the pyridyl group),
(6) m is 2,
(7) R⁴ is hydrogen,
25 (8) R⁴ is C1-C2 alkyl,
(9) R⁵ is C1-C2 alkyl, phenyl (which may have 1 to 3 substituent(s)), pyridyl (which may have 1 to 3 substituent(s))
(10) R⁵ is pyridyl (which may have 1 to 3 substituent(s)),
30 (11) n is 2,

(12) substituent(s) is(are) cyano.

The compound of the formula (I) of the present invention can be prepared according to the following
 5 process A-1 to A-3.

[Process A-1]



10 In the above formulae, R^1 , R^2 and X represent the same meanings as defined above. And "Hal" represents halogen atom, especially, chlorine or bromine atom.

Process A-1 is the process for preparing the Compound (I) from amine Compound (VI) and halogen Compound (VII)
 15 in solvent, preferably in the presence of base.

Compound (VI) and (VII) may be purchased if it is commercial, or synthesized according to Processes C and B, respectively, mentioned after or other general methods obvious to the person skilled in the organic chemistry
 20 from commercial compounds.

The solvent employable in this process is not particularly limited so long as it is inactive in this reaction and may include ethers such as diisopropyl ether, tetrahydrofuran, dioxane, preferably tetrahydrofuran.

25 The base employable in this process for making basic condition is not particularly limited so long as it accelerates this reaction and may include organic amines

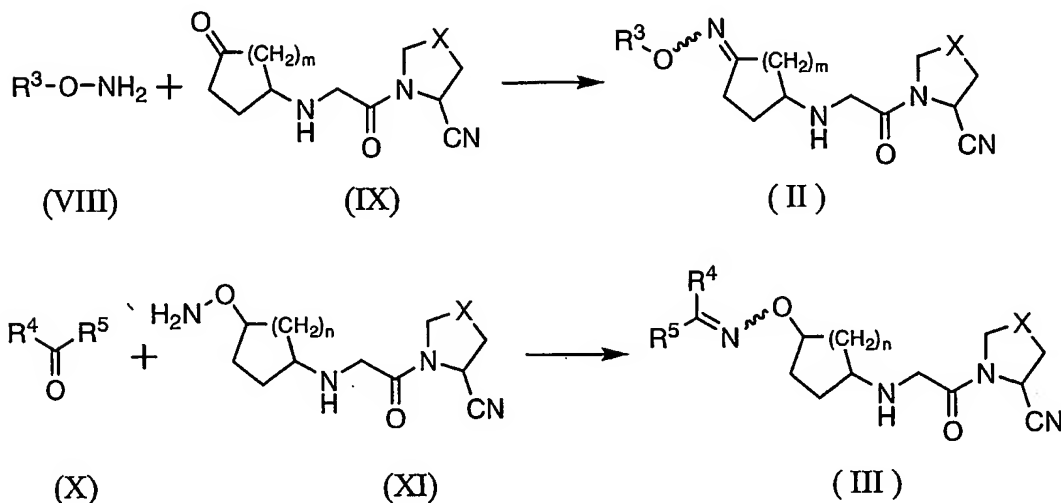
such as triethylamine, tributylamine, diisopropylethylamine; alkali metal hydrogencarbonates such as lithium hydrogencarbonate, sodium hydrogencarbonate and potassium hydrogencarbonate; 5 alkali metal carbonates such as lithium carbonate, sodium carbonate and potassium carbonate; alkaline earth metal carbonates such as magnesium carbonate and calcium carbonate; alkali metal hydroxides such as lithium hydroxide, sodium hydroxide and potassium hydroxide, 10 preferably organic amine, especially triethylamine.

This process is generally carried out by adding the solution of Compound (VII) preferably dropwise to the solution of Compound (VI). The temperature at that time varies depending on the starting material, the solvent, 15 etc., but it is usually from -10°C to 20°C . And after adding, the temperature is preferably raised to room temperature.

The reaction time after the adding varies depending on the starting material, the solvent, etc., but it is 20 usually from 1hr to 24hrs, preferably from 1hr to 15hrs.

After the reaction, the mixture is quenched by aqueous solvent such as water, saturated NH_4Cl , hydrochloric acid, etc, and extracted with organic solvent insoluble with water such as ethyl acetate, CHCl_3 , etc. If the target 25 compound is in aqueous layer, base such as NaHCO_3 is added. Preferably, the organic layer is washed with water or the like, dried over anhydrous MgSO_4 or Na_2SO_4 , evaporated in vacuo, and the target compound is purified by the conventional method such as thin layer chromatography, 30 silica gel column chromatography, etc.

Compound (II) and (III) included in the scope of Compound (I) can be synthesized by following Process A-2.
[Process A-2]



In the above formulae, R^3 to R^5 , X, m and n represent the same meanings as defined above.

Process A-2 is the process for preparing the Compound (II) and (III) by forming oximino group from aminoxy compound and carbonyl compound in solvent. And Compound (II) and (III) are the Compound (I), of which the definition R^1 is not hydrogen or lower alkyl.

Starting Compound (VIII) to (XI) may be purchased if they are commercial, or synthesized according to general methods from commercial compounds. For example, aminoxy group of Compound (VIII) and (XI) can be synthesized by application of Process D described later.

The solvent employable in this process is not particularly limited so long as it is inactive in this reaction and may include alcohols such as methanol,

ethanol, isopropanol, preferably methanol. But, in case that Compound (X) may be used as solvent, other solvent is not necessarily needed.

5 This process is generally carried out by adding the solution of carbonyl compound (Compound (IX) or (X)) to the solution of aminoxy compound (Compound (VIII) or (XI)). The temperature at that time varies depending on the starting material, the solvent, etc., but it is usually from 10°C to 50°C, preferably room temperature.

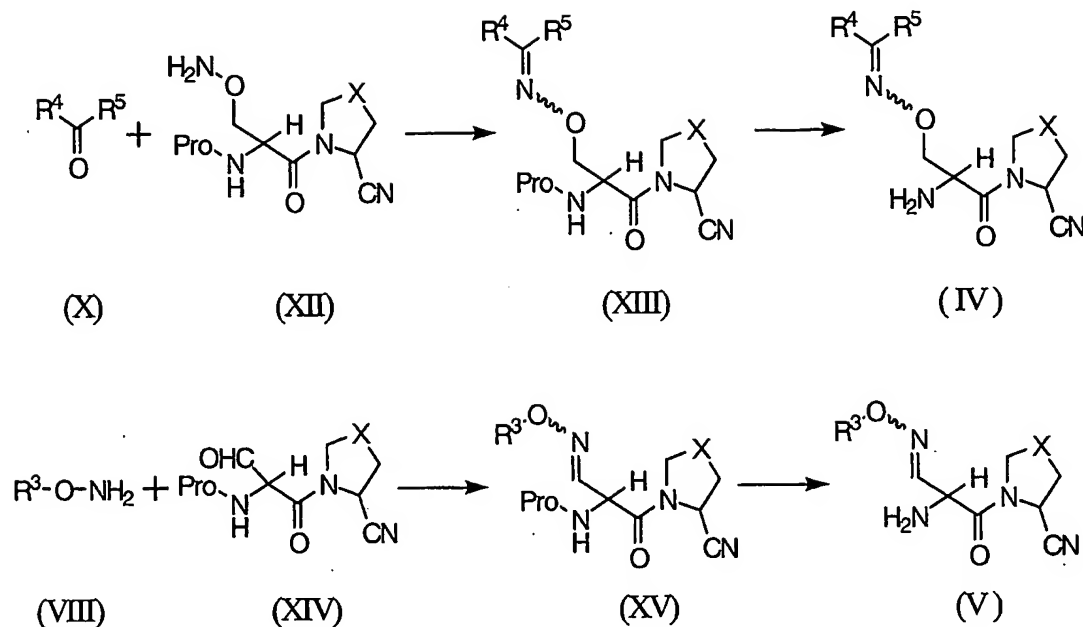
10 The reaction time after the adding varies depending on the starting material, the solvent, etc., but it is usually from 30min to 12hrs, preferably from 1hr to 5hrs.

After the reaction, the solvent is removed in vacuo, and the residue is dissolved in organic solvent insoluble with water such as ethyl acetate, CHCl_3 , etc. The solution
15 is washed with aqueous solvent such as brine, and dried over anhydrous MgSO_4 or Na_2SO_4 , evaporated in vacuo. The target compound may be purified by the conventional method such as silica gel column chromatography, etc.

20

Compound (IV) and (V) included in the scope of Compound (I) can be synthesized by following Process A-3.

[Process A-3]



In the above formulae, R³ to R⁵ and X represent the same meanings as defined above. And "Pro" means the protective group of amino group.

Process A-3 is the process for preparing the Compound (IV) and (V) by forming oxyimino group from aminoxy compound and carbonyl compound in solvent similar to Process A-2, then deprotecting step. And Compound (IV) and (V) are the Compound (I), of which the definition R² is not hydrogen.

In the first step (forming oxyimino group), the same procedure as that in Process A-2 may be adopted in this process, because forming of oxyimino group is common to both of Process.

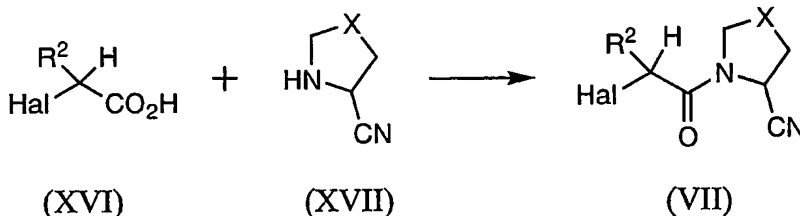
In the second step (deprotecting), concerning the kind of protective group ("Pro") and cleavage reaction of the protective group, [PROTECTIVE GROUPS IN ORGANIC SYNTHESIS Second Edition] T.W.Green and P.G.M.Wuts, John

Wiley & Sons, INC. may be referred.

For example, when "Pro" group is carbamate group such as tert-butoxycarbonyl group, the cleavage reaction is carried out in the acidic condition in solvent by acid
 5 such as hydrochloric acid, trifluoroacetic acid, or the like. After the deprotection, excess acid and solvent are removed in vacuo, and the target compound may be purified by triturating, which uses appropriate solvent such as diethylether.

10

Compound (VII), which is the starting compound of Process A-1, can be synthesized by following Process B.
 [Process B]



15

In the above formulae, R², X and "Hal" represent the same meanings as defined above.

Process B is the process for preparing the Compound (VII) by forming amide bond in solvent in the presence of dehydrating agent, base and catalyst.

20

Compound (XVI) and (XVII) may be purchased if it is commercial, or synthesized according to general methods obvious to the person skilled in the organic chemistry from commercial compounds.

The solvent employable in this process is not particularly limited so long as it is inactive in this
 25 reaction and may include amides such as dimethylformamide and dimethylacetamide.

The dehydrating agent employable in this process is not particularly limited so long as it accelerate forming amide bond and may include carbodiimide compounds such as dicyclohexylcarbodiimide (DCC),
5 diisopropylcarbodiimide, water solvable carbodiimide (WSCD), preferably WSCD.

The base employable in this process is not particularly limited so long as it accelerates this process and may include organic amines such as
10 triethylamine, tributylamine, diisopropylethylamine (DIEA), preferably DIEA.

The catalyst employable in this process is not particularly limited so long as it can mainly make the carboxyl group of Compound (XVI) active and may include
15 1-hydroxybenzotriazole (HOBT).

This process is generally carried out by adding the dehydrating agent and base to the solution of Compound (XVI), Compound (XVII) and catalyst. The temperature at that time varies depending on the starting material, the
20 solvent, etc., but it is usually from -10°C to 30°C, preferably from 0°C to 10°C. And after adding, the temperature is preferably raised to room temperature.

The reaction time after the adding varies depending on the starting material, the solvent, etc., but it is
25 usually from 2hrs to 24hrs.

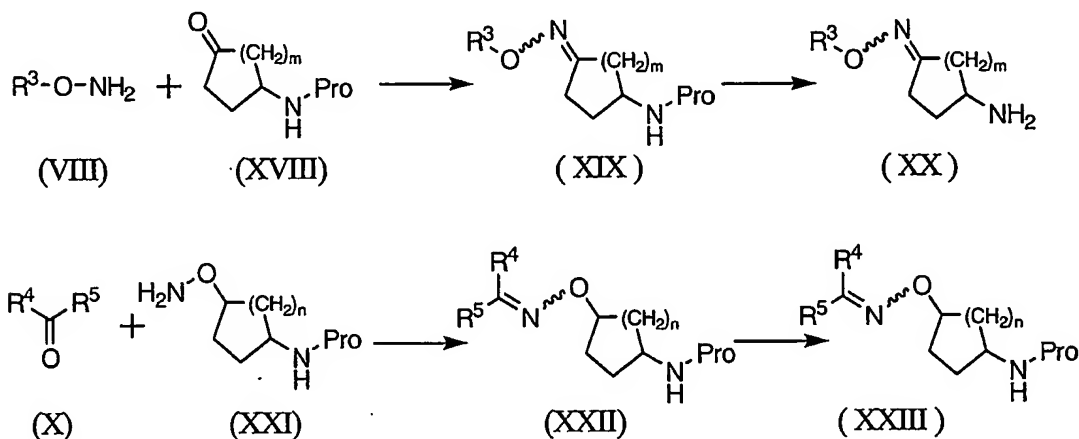
After the reaction, the solvent is removed in vacuo, the target compound may be purified by the conventional method such as silica gel column chromatography, etc.

30 Compound (XX) and (XXIII), both of which are the

starting Compound (VI) of Process A-1, can be synthesized by following Process C. Provided that Compound (XX) and (XXIII) are the Compound (VI), of which the R¹ is not hydrogen or lower alkyl.

- 5 Incidentally, in case that R¹ of Compound (VI) is hydrogen or lower alkyl, its structure is very simple. So, the compound may be purchasable.

[Process C]



10

In the above formulae, R³ to R⁵, m, n and "Pro" represent the same meanings as defined above.

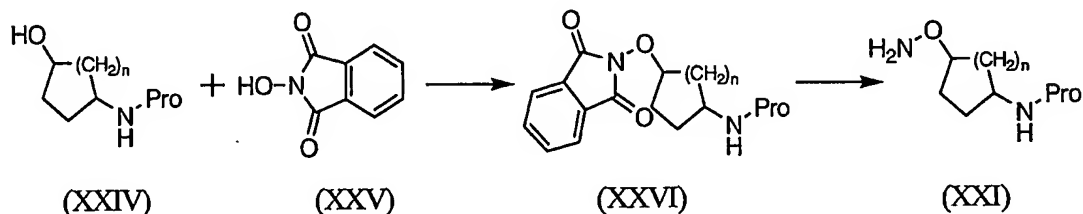
- Process C is the process for preparing the Compound (XX) and (XXIII) by forming oxyimino group from amino compound and carbonyl compound in solvent.
- 15

And the same procedure as that in Process A-2 may be adopted in this process, because forming of oxyimino group is common to both of Process.

- 20 The compound having aminooxy group such as Compound (VIII) and (XI) (the starting compound of Process A-2, and the like), Compound (XII) (the starting compound of Process A-3), Compound (XXI) (the starting compound of

Process C), can be synthesized by following Process D. But, in the following reaction formula, the synthesis process of Compound (XXI) is shown as representative example.

5 [Process D]



In the above formulae, n and "Pro" represent the same meanings as defined above.

Process D is the process for preparing the compound having aminooxy group by functional group transformation from hydroxy group.

In this process, first, to the solution of the compound having hydroxy group (Compound (XXIV)), Compound (XXV) and triphenylphosphine was added diisopropyl azodicarboxylate (DIAD).

The compound having hydroxy group may be purchased if it is commercial, or synthesized according to general methods obvious to the person skilled in the organic chemistry from commercial compounds. And Compound (XXV) may be purchasable.

The solvent employable in this process is not particularly limited so long as it is inactive in this reaction and may include ethers such as diisopropyl ether, tetrahydrofuran, dioxane, preferably tetrahydrofuran.

The temperature when adding the dehydrating agent (DIAD) varies depending on the starting material, the solvent, etc., but it is usually from -10°C to 50°C,

preferably from 0°C to 30°C.

The reaction time after the adding varies depending on the starting material, the solvent, etc., but it is usually from 30min to 12hrs.

5 After the reaction, to the solution water, then extracted with organic solvent insoluble with water such as ethyl acetate, CHCl_3 , etc. The organic layer is dried over anhydrous MgSO_4 or Na_2SO_4 , evaporated in vacuo. And by adding the solvent to dissolve the target compound but
10 not to able dissolve triphenylphosphonate, the target compound (phthalimide derivative) may be obtained. The phthalimide derivative may be used in next step without further purification.

Then, phthalimide derivative (Compound (XXVI)) was
15 decomposed by hydrazine in solvent. That is, to the solution of phthalimide derivative, hydrazine is added.

The solvent employable in this process is not particularly limited so long as it is inactive in this reaction and may include alcohols such as methanol,
20 ethanol, isopropanol, preferably ethanol.

The temperature at that time varies depending on the starting material, the solvent, etc., but it is usually from 50°C to 150°C, preferably from 70°C to 130°C.

The reaction time after the adding varies depending
25 on the starting material, the solvent, etc., but it is usually from 5min to 5hrs.

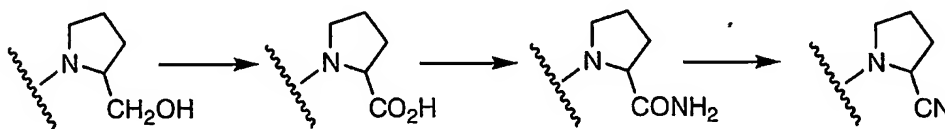
After the reaction, the resulting precipitate is filtered off and washed with solvent, and filtrate is concentrated. Then, to the residue, aqueous solvent and
30 organic solvent insoluble with water such as ethyl acetate

are added, aqueous layer is basified and extracted. The organic layer is dried over anhydrous MgSO_4 or Na_2SO_4 , evaporated in vacuo to obtain the target compound. If necessary, further purification may be carried out.

5

In the above processes, functional group transformation may be carried out on cue so long as the other sites of the compounds are not affected. In the following reaction formula, the functional group transformation of the pyrrolidine ring is shown as representative example.

[Process E]



Process E shows the representative example of functional group transformation. Accordingly, other reactions of functional group transformation may be carried out.

Above processes, all starting materials and product compounds may be salts. The compounds of above processes can be converted to salt according to a conventional method. For example of making hydrochloride or trifluoroacetate, to the compound, the solution of hydrochloride such as 4N hydrochloride/dioxane or trifluoroacetic acid is added, then the solvent and excess acid is removed and the residue is triturated appropriate solvent such as diethylether.

For therapeutic purpose, the Compound (I) and a

pharmaceutically acceptable salt thereof of the present invention can be used in a form of pharmaceutical preparation containing one of said compounds as an active ingredient, in admixture with a pharmaceutically acceptable carrier such as an organic or inorganic solid or liquid excipient suitable for oral, parenteral or external administration. The pharmaceutical preparations may be capsules, tablets, dragees, granules, inhalant, suppositories, solution, lotion, suspension, emulsion, ointment, gel, cream, or the like. If desired, there may be included in these preparations, auxiliary substances, stabilizing agents, wetting or emulsifying agents, buffers and other commonly used additives.

While the dosage of therapeutically effective amount of the Compound (I) will vary depending upon the age and condition of each individual patient, an average single dose of about 0.01 mg, 0.1 mg, 1 mg, 10 mg, 50 mg, 100 mg, 250 mg, 500 mg and 1000 mg of the Compound (I) may be effective for treating the above-mentioned diseases. In general, amounts between 0.01 mg/body and about 1,000 mg/body may be administered per day.

Best Mode for Carrying Out the Invention

The patents, patent applications and publications cited herein are incorporated by reference.

5 The following Examples are given only for the purpose of illustrating the present invention in more detail.

Abbreviations used in this application are as follows,
TLC : thin layer chromatography.

10

Example 1-1

tert-Butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-
2-oxo-1-[[[(3-pyridinylmethylidene)amino]oxy]methyl]e
thylcarbamate

15

To a solution of tert-butyl
(1S)-1-[(aminooxy)methyl]-2-[(2S)-2-carbamoyl-1-pyrro
lidinyl]-2-oxoethylcarbamate (120mg) in methanol (5ml)
was added nicotinaldehyde (0.036ml) at room temperature.
20 The reaction mixture was set aside at room temperature
for 2hrs.

After removal of the solvent in vacuo, the residue
was dissolved in ethyl acetate (10ml), washed with
brine(5ml), and dried over MgSO₄. After removal of the
25 solvent in vacuo, the residue was purified with silica
gel TLC (0.5mm X 2, CHCl₃:methanol=20:1) to give the target
compound as a colorless oily amorphous (120mg, 78%).

¹H NMR(CDCl₃) : 1.45(s, 9H), 1.90-2.40(m, 4H), 3.60-3.77(m,
30 2H), 4.34-4.40(m, 2H), 4.64-4.69(m, 1H), 4.85-4.94(m 1H),

5.21-5.28(m, 1H), 5.37-5.44(m, 1H), 6.68(br-s, 1H),
7.28-7.36(m, 1H), 7.90-7.95(m, 1H), 8.10(s, 1H),
8.62-8.65(m, 1H), 8.74(s, 1H)

MS : 406.17 (ES+)

5

Example 1-2

tert-Butyl (1S)-2-[(2S)-2-cyano-1-pyrrolidinyl]-2-oxo-1-[[[(3-pyridinylmethylidene)amino]oxy]methyl]ethylcarbamate

10

To a slurry of tert-butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-2-oxo-1-[[[(3-pyridinylmethylidene)amino]oxy]methyl]ethylcarbamate (120mg) obtained by Example 1-1 in tetrahydrofuran (3ml) were added triethylamine (0.1ml) and trifluoroacetic anhydride (0.084ml) dropwise at 5°C (clearly solved). The reaction mixture was stirred at 5°C for 30min.

Sat. NaHCO₃ was added and extracted with ethyl acetate. The organic layer was dried over MgSO₄. After removal of the solvent, the residue was purified with TLC (0.5mm X 1, CHCl₃:methanol=20:1) to give the target compound as a colorless oily amorphous (75mg, 65.4%).

¹H NMR(CDCl₃) : 1.43(s, 9H), 2.06-2.31(m, 4H), 3.65-3.81(m, 2H), 4.35-4.40(m, 2H), 4.76-4.85(m, 2H), 5.32-5.40(m, 1H), 7.28-7.36(m, 1H), 7.94-8.00(m, 1H), 8.13(s, 1H), 8.60-8.65(m, 1H), 8.74(s, 1H)

MS : 388.14 (ES+)

30 Example 1-3

(2S)-1-[(2S)-2-Amino-3-[[[(3-pyridinylmethylidene)amino]oxy]propanoyl]-2-cyanopyrrolidine trifluoroacetate

tert-Butyl (1S)-2-[(2S)-2-cyano-1-pyrrolidinyl]-
5 2-oxo-1-[[[(3-pyridinylmethylidene)amino]oxy]methyl]ethylcarbamate (70mg) obtained by Example 1-2 dissolved in trifluoroacetic acid (2ml) at room temperature, and the reaction mixture was stirred at room temperature for 30min.

10 After removal of the solvent in vacuo, the residue was triturated with diethylether to give the target compound as a pale yellow solid (45mg, 62.1%).

¹H NMR(DMSO-d₆) : 1.95-2.33(m, 4H), 3.64-3.79(m, 2H),
15 4.44-4.64(m, 3H), 4.77-4.87(m, 1H), 7.50-7.58(m, 1H),
8.14-8.21(m, 1H), 8.40(s, 1H), 8.41-8.89(m, 4H)
MS : 288.12(ES+), free 287.32

Example 2-1

20 cis-tert-Butyl 4-[(1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl)oxy]cyclohexylcarbamate

To a solution of trans-tert-butyl
4-hydroxycyclohexylcarbamate (3g),
25 2-hydroxy-1H-isoindole-1,3(2H)-dione (2.27g), and
triphenylphosphine (3.84g) in tetrahydrofuran (30ml) was
added diisopropyl azodicarboxylate (DIAD, 3.02ml)
dropwise at 5°C. The reaction mixture was stirred at room
temperature for 2hrs.

30 Then, water (10ml) was added and extracted with ethyl

acetate (50ml). The organic layer was dried over MgSO_4 and filtered. After removal of the solvent in vacuo, the residue was triturated with isopropylether, the resulting precipitate (triphenylphosphonate) was filtered off.
5 The filtrate was concentrated to give the yellow oil (5.0g, 99.6%). Further purification was not attempted.

Example 2-2

cis-tert-Butyl 4-(aminooxy)cyclohexylcarbamate

10

To a solution of the crude cis-tert-butyl 4-[(1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl)oxy]cyclohexylcarbamate (5g) obtained by Example 2-1 in ethanol (50ml) was added hydrazine hydrate (7ml) at 120°C (bath
15 temp.). After 10min, the reaction mixture gave white precipitate (phthalhydrazine).

The resulting precipitate was filtered off, and the filtrate was concentrated. The residue was dissolved in 1N HCl (40ml) and ethyl acetate (40ml). The aqueous layer
20 was basified with NaHCO_3 , and saturated with NaCl, and extracted with ethyl acetate (40ml). The organic layer was dried over MgSO_4 . After removal of the solvent in vacuo, the residue was crystallized from isopropylether and collected by filtration to give the target compound
25 as a white powder (1.16g, 36.3%).

^1H NMR(CDCl_3) : 1.44(s, 9H), 1.45-1.91(m, 8H), 3.50(br-s, 1H), 3.64(br-s, 1H), 4.48(br-s, 1H), 5.23(s, 2H)
MS : 231.17 (ES+)

30

Example 2-3

cis-tert-Butyl 4-[[[(3-pyridinylmethylidene)amino]-oxy]cyclohexylcarbamate

5 To a solution of cis-tert-butyl 4-(aminooxy)cyclohexylcarbamate (200mg) obtained by Example 2-2 in methanol (4ml) was added nicotinaldehyde (93mg) at room temperature. The reaction mixture was set aside at room temperature for 2hrs.

10 After removal of the solvent in vacuo, the residue was dissolved in ethyl acetate, washed with brine, and dried over MgSO₄. After removal of the solvent in vacuo, the target compound was obtained as a colorless oily amorphous (200mg, 72.1%). Further purification was not
15 attempted.

¹H NMR(CDCl₃) : 1.45(s, 9H), 1.46-2.06(m, 8H), 3.55(br-s, 1H), 4.33(br-s, 1H), 4.50(br-s, 1H), 7.28-7.32(m, 1H), 7.95-7.98(m, 1H), 8.10(s, 1H), 8.57-8.60(m, 1H), 8.73(s, 1H)
20

MS : 320.18(ES+)

Example 2-4

cis-Nicotinaldehyde O-(4-aminocyclohexyl)oxime

25

cis-tert-Butyl 4-[[[(3-pyridinylmethylidene)-aminooxy]cyclohexylcarbamate (200mg) obtained by Example 2-3 was dissolved in small amount of methanol, and 4N HCl-dioxane (5ml) was added at room temperature.

30 After 30min, the solvent was removed in vacuo, the

residue was basified with sat. NaHCO₃ and extracted with CHCl₃ (5 times). The organic layer was dried over MgSO₄ and filtered. After removal of the solvent in vacuo, the target compound was obtained as colorless oil (120mg, 87.4%).

¹H NMR(CDCl₃) : 1.46-2.15(m, 8H), 2.16-2.37(br-s, 2H), 2.80-2.90(m, 1H), 4.34-4.40(m, 1H), 7.30-7.32(m, 1H), 7.93-7.97(m, 1H), 8.12(s, 1H), 8.57-8.60(m, 1H), 8.74(s, 1H)
MS : 220.14 (ES+)

Example 2-5

cis-(2S)-1-[[[4-[(3-Pyridinylmethylidene)amino]oxy]cyclohexyl]amino]acetyl]-2-cyanopyrrolidine dihydrochloride

To a solution of cis-nicotinaldehyde O-(4-aminocyclohexyl)oxime (120mg) obtained by Example 2-4 and triethylamine (0.077ml) in tetrahydrofuran (4ml) was added a solution of (2S)-1-bromoacetyl-2-cyanopyrrolidine (60mg) in tetrahydrofuran (1ml) dropwise at 5°C.

After 2hrs, the reaction mixture was diluted with sat. NH₄Cl and extracted with ethyl acetate. The organic layer was dried over MgSO₄ and filtered. After removal of the solvent in vacuo, the residue was purified with TLC (0.5mm X 1, CHCl₃:methanol=10:1).

After removal of the solvent in vacuo, the oily amorphous was dissolved in methanol and 4N HCl-dioxane.

After removal of the solvent, the residue was triturated with diethylether and the solvent was removed in vacuo. The target compound was obtained as a white powder (50mg, 42.2%).

5

^1H NMR(DMSO- d_6) : 1.60-2.22(m, 12H), 3.13(br-s, 1H), 3.45-3.74(m, 2H), 3.90-4.15(m, 2H), 4.41(br-s, 1H), 4.85(m, 1H), 7.89-7.95(m, 1H), 8.47(s, 1H), 8.57-8.60(m, 1H), 8.86(m, 1H), 9.06(s, 1H), 9.20(br-s, 1H)

10 MS : 356.13(ES+), free 355.43

Example 3-1

cis-tert-Butyl 4-[[[(1-methylethylidene)amino]oxy]-cyclohexylcarbamate

15

To a solution of cis-tert-butyl 4-(aminooxy)cyclohexylcarbamate (400mg) in methanol (2ml) was added acetone (10ml) at room temperature. The reaction mixture was set aside at room temperature for 20 2hrs.

After removal of the solvent in vacuo, the residue was dissolved in ethyl acetate, washed with brine, and dried over MgSO_4 . After removal of the solvent in vacuo, the residue was crystallized from n-hexane to give the 25 target compound as a white solid (460mg, 98%).

^1H NMR(CDCl_3) : 1.40-1.95(m, 8H), 1.44(s, 9H), 1.86(s, 3H), 1.87(s, 3H), 3.60-3.67(m, 1H), 4.13-4.19(m, 1H)
MS : 271.21(ES+)

30

Example 3-2

Acetone O-(4-aminocyclohexyl)oxime

tert-Butyl 4-[[[(1-methylethylidene)amino]oxy]-
5 cyclohexylcarbamate (460mg) obtained by Example 3-1 was
dissolved in small amount of methanol, and 4N HCl-dioxane
(5ml) was added at room temperature.

After 30min, the solvent was removed in vacuo, the
residue was basified with sat. NaHCO₃ and extracted with
10 CHCl₃ (5 times). The organic layer was dried over MgSO₄
and filtered. After removal of the solvent in vacuo, the
residual solid was triturated with isopropylether and
collected by filtration to give the target compound as
a white powder (200mg, 69%).

15

¹H NMR(CDCl₃) : 1.45-2.14(m, 8H), 1.87(s, 3H), 1.91(s,
3H), 3.15-3.25(m, 1H), 4.20-4.25(m, 1H)

MS : 171.04(ES+)

20 Example 3-3

cis-(2S)-1-[[[4-[[[(1-Methylethylidene)amino]oxy]cyclo
hexyl]amino]acetyl]-2-cyanopyrrolidine hydrochloride

To a solution of cis-acetone
25 O-(4-aminocyclohexyl)oxime (200mg) obtained by Example
3-2 and triethylamine (0.17ml) in tetrahydrofuran (4ml)
was added a solution of
(2S)-1-bromoacetyl-2-cyanopyrrolidine (130mg) in
tetrahydrofuran (1ml) dropwise at 5°C.

30 After 12hrs, the solvent was removed in vacuo, the

residue was treated with diethylether and 1N HCl. The aqueous layer was separated, basified with NaHCO₃, and extracted with ethyl acetate. The organic layer was dried over MgSO₄ and filtered. After removal of the solvent in vacuo, the residue was purified with TLC (0.5mm X 1, CHCl₃:methanol=20:1). The residue was dissolved in small amount of methanol and 4N HCl-dioxane was added. After removal of the solvent in vacuo, the target compound was obtained as a white solid (25mg, 12.2%).

10

¹H NMR(DMSO-d₆) : 1.40-2.23(m, 12H), 1.82(s, 3H), 1.86(s, 3H), 3.38-4.90(m, 7H)

MS : 307.20(ES+), free 306.40

15 Example 4-1

cis-tert-Butyl 4-[[[(2-pyridinylmethylidene)amino]oxy]cyclohexylcarbamate

To a solution of cis-tert-butyl 4-(aminooxy)cyclohexylcarbamate (200mg) in methanol (4ml) was added 2-pyridinecarbaldehyde (0.083ml) at room temperature. The reaction mixture was set aside at room temperature for 2hrs.

After removal of the solvent in vacuo, the target compound was obtained as a yellow oil (260mg, 93.7%). Further purification was not attempted.

Example 4-2

cis-2-Pyridinecarbaldehyde O-(4-aminocyclohexyl)oxime

30

cis-tert-Butyl 4-[[[(2-pyridinylmethylidene)-amino]oxy]cyclohexyl]carbamate (260mg) obtained by Example 4-1 was dissolved in small amount of methanol, and 4N HCl-dioxane (5ml) was added at room temperature.

5 After 30min, the solvent was removed in vacuo, the residue was basified with sat. NaHCO₃ and extracted with CHCl₃ (5 times). The organic layer was dried over MgSO₄ and filtered. After removal of the solvent in vacuo, the target compound was obtained as colorless oil (180mg, 10 about 100%).

¹H NMR(CDCl₃) : 1.55-2.27(m, 8H), 3.16-3.27(m, 1H), 3.71(s, 2H), 4.40-4.45(m, 1H), 7.23-7.30(m, 1H), 7.64-7.75(m, 2H), 8.22(s, 1H), 8.61-8.62(m, 1H)

15 MS : 220.18(ES+)

Example 4-3

cis-(2S)-1-[[[4-[[[(2-Pyridinylmethylidene)amino]oxy]cyclohexyl]amino]acetyl]-2-cyanopyrrolidine
20 dihydrochloride

To a solution of cis-2-pyridinecarbaldehyde O-(4-aminocyclohexyl)oxime (180mg) obtained by Example 4-2 and triethylamine(0.116ml) in tetrahydrofuran (2ml)
25 was added a solution of (2S)-1-bromoacetyl-2-cyanopyrrolidine (90mg) in tetrahydrofuran (1ml) dropwise at 5°C.

After 2hrs stirring at room temperature, the reaction mixture was diluted with sat. NH₄Cl and extracted with ethyl
30 acetate. The organic layer was dried over MgSO₄ and

filtered. After removal of the solvent in vacuo, the residue was purified with TLC (0.5mm, CHCl₃:methanol=10:1). After removal of the solvent in vacuo, the residual oily amorphous was dissolved in methanol and 4N HCl-dioxane. After removal of the solvent, the residue was triturated with diethylether and the solvent was removed in vacuo, the target compound was obtained as a white powder (50mg, 28.2%).

¹H NMR(DMSO-d₆) : 1.56-2.25(m, 12H), 3.17(br-s, 1H), 3.44-3.70(m, 2H), 3.97-4.40(m, 3H), 4.80-4.86(m, 1H), 7.60-7.75(m, 1H), 7.93-7.98(m, 1H), 8.05-8.13(m, 1H), 8.36(s, 1H), 8.68-8.70(m, 1H), 9.13(br-s, 1H)
MS : 356.24(ES+), free 355.43

15

Example 5-1

tert-Butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-1-hydroxymethyl-2-oxoethylcarbamate

To a solution of (2S)-2-[(tert-butoxycarbonyl)amino]-3-hydroxypropanoic acid (5g), (2S)-2-pyrrolidinecarboxamide (2.78g), and 1-hydroxybenzotriazole hydrate (HOBt-H₂O, 4.47g) in N,N-dimethylformamide (DMF, 50ml) were added 1-[3-(dimethylamino)propyl]-3-ethylcarbodiimide hydrochloride (WSCD-HCl, 5.14g) and diisopropylethylamine (DIEA, 5.1ml) at 5°C. The reaction mixture was stirred at room temperature overnight.

The solvent was removed in vacuo, and the resulting residue was purified with silica gel column chromatography.

(SiO₂, 200ml, CHCl₃:methanol=10:1). After removal of the solvent in vacuo, the residual oil was triturated with diethylether and solidified, washed with diethylether to give the target compound as a white powder (5g, 68.1%).

5

¹H NMR(CDCl₃) : 1.43(s, 9H), 1.90-2.22(m, 4H), 3.56-3.96(m, 4H), 4.25-4.70(m, 3H), 5.75(m, 1H), 6.31(s, 1H), 6.94(s, 1H)

MS : 302.16 (ES+)

10

Example 5-2

tert-Butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-1-[[[(1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl)oxy]methyl]-2-oxoethylcarbamate

15

To a solution of tert-butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-1-hydroxymethyl-2-oxoethylcarbamate (2g) obtained by Example 5-1, 2-hydroxy-1H-isoindole-1,3(2H)-dione (1.08g) and triphenylphosphine (1.8g) in tetrahydrofuran (20ml) was added diisopropyl azodicarboxylate (DIAD, 1.44ml) dropwise at 5°C. The reaction mixture was stirred at 5°C for 1hr.

The resulting gel was collected by filtration, washed with tetrahydrofuran. The collected solid was used for the next step without further purification (3g, about 100%).

Example 5-3

tert-Butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-

1-[(aminooxy)methyl]-2-oxoethylcarbamate

To a solution of tert-butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-1-[[[(1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl)oxy]methyl]-2-oxoethylcarbamate (900mg) obtained by Example 5-2 in ethanol (10ml) was added hydrazine hydrate (1ml) at 120°C (bath temp.). After 10min, the reaction mixture gave white precipitate (phthalhydrazine).

10 The resulting precipitate was filtered off, and the filtrate was concentrated. The residue was dissolved in 1NHCl and ethyl acetate. The aqueous layer was separated, basified with NaHCO₃ carefully, saturated with NaCl, and extracted with CHCl₃. The organic layer was dried over 15 MgSO₄ and filtered. After removal of the solvent in vacuo, the target compound was obtained as a colorless oily amorphous (800mg, 37.6%).

¹H NMR(CDCl₃) : 1.45(s, 9H), 1.62-1.71(m, 2H), 2.00-2.07(m, 20 2H), 2.32-2.37(m, 1H), 3.67-3.91(m, 4H), 4.63-4.78(m, 2H), 5.36-5.70(m, 4H)
MS : 317.20(ES+)

Example 5-4

25 tert-Butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-2-oxo-1-[[[(2-pyridinylmethylidene)amino]oxy]methyl]ethylcarbamate

To a solution of tert-butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-1-[(aminooxy

)methyl]-2-oxoethylcarbamate (200mg) obtained by Example 5-3 in methanol (4ml) was added 2-pyridinecarbaldehyde (0.06ml) at room temperature. The reaction mixture was set aside at room temperature for 2hrs.

5 After TLC check, the solvent was removed in vacuo, the residue was dissolved in ethyl acetate, washed with brine, and dried over MgSO_4 . After removal of the solvent in vacuo, the residue was purified with TLC (0.5mm X 2, CHCl_3 :methanol=20:1) to give the target compound as a
10 colorless oily amorphous (120mg, 46.8%).

^1H NMR(CDCl_3) : 1.44(s, 9H), 1.91-2.34(m, 5H), 3.73(m, 1H), 4.40-4.44(m, 2H), 4.65(m, 1H), 4.90(m, 1H), 5.49(m, 2H), 6.73(br-s, 1H), 7.27-7.30(m, 1H), 7.71-7.75(m, 2H),
15 8.16(s, 1H), 8.60-8.62(m, 1H)

MS : 406.21(ES+)

Example 5-5

tert-Butyl (1S)-2-[(2S)-2-cyano-1-pyrrolidinyl]-2-
20 oxo-1-[[[(2-pyridinylmethylidene)amino]oxy]methyl]ethylcarbamate

To a slurry of tert-butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-2-oxo-1-[[[(2-pyridinylmethylidene)amino]oxy]methyl]ethylcarbamate
25 e (120mg) obtained by Example 5-4 in tetrahydrofuran (2.4ml) were added triethylamine (0.13ml) and trifluoroacetic anhydride (TFAA, 0.084ml) dropwise at 5°C.

30 The reaction mixture was stirred at 5°C for 30min,

then sat. NaHCO₃ was added, and extracted with ethyl acetate. The organic layer was dried over MgSO₄. After removal of the solvent, the residue was purified with TLC (0.5mm X 1, CHCl₃:methanol=20:1) to give the target compound
5 as a colorless oily amorphous (90mg, 78.5%).

¹H NMR(CDCl₃) : 1.42(s, 9H), 2.17-2.30(m, 4H), 3.65-3.84(m, 2H), 4.37-4.42(m, 2H), 4.75-4.87(m, 2H), 5.30-5.37(m, 1H), 7.27-7.30(m, 1H), 7.65-7.85(m, 2H), 8.20(s, 1H),
10 8.59-8.62(m, 1H)

MS : 388.16(ES+)

Example 5-6

(2S)-1-[(2S)-2-Amino-3-[[(2-pyridinylmethylidene) amino
15 o]oxy]propanoyl]-2-cyanopyrrolidine dihydrochloride

tert-Butyl (1S)-2-[(2S)-2-cyano-1-pyrrolidinyl]-2-oxo-1-[[[(2-pyridinylmethylidene)amino]oxy]methyl]ethylcarbamate (70mg) obtained by Example 5-5 was dissolved
20 in trifluoroacetic acid (TFA, 2ml) at room temperature, and the reaction mixture was stirred at room temperature for 30min.

After removal of the solvent in vacuo, the residual oil was dissolved in sat. NaHCO₃ and extracted with CHCl₃.
25 The organic layer was dried over MgSO₄ and filtered. After removal of the solvent in vacuo, the residue was treated with 4NHCl-dioxane. After removal of the solvent in vacuo, the residue was triturated with diethylether, and the solvent was removed in vacuo. The target compound was
30 obtained as a pale yellow amorphous powder (35mg, 53.8%).

¹H NMR(DMSO-d₆) : 1.95-2.32(m, 4H), 3.45-3.80(m, 3H),
4.51-4.62(m, 2H), 4.80-4.86(m, 1H), 7.50-7.55(m, 1H),
7.93(m, 2H), 8.27(s, 1H), 8.55-8.65(br-s, 2H),
5 8.65-8.70(m, 1H)
MS : 288.16(ES+), free 287.32

Example 6-1

tert-Butyl (1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-
10 1-[[[(1-methylethylidene)amino]oxy]methyl]-2-oxoethyl
carbamate

tert-Butyl (1S)-2-[(2S)-2-carbamoyl-1-
pyrrolidinyl]-1-[(aminooxy)methyl]-2-oxoethylcarbamate
15 e (300mg) was dissolved in acetone (10ml) at room
temperature. The reaction mixture was set aside at room
temperature for 10min.

The solvent was removed in vacuo, the residue was
dissolved in ethyl acetate, washed with 1N HCl, sat. NaHCO₃,
20 brine, and dried over MgSO₄. After removal of the solvent
in vacuo, the residue was crystallized from
diethylether-isopropylether to give the target compound
as a white powder (150mg, 44.4%).

25 ¹H NMR(CDCl₃) : 1.44(s, 9H), 1.82(s, 3H), 1.84(s, 3H),
1.93-2.40(m, 4H), 3.61-3.73(m, 2H), 4.15-4.25(m, 2H),
4.63-4.70(m, 1H), 4.73-4.82(m, 1H), 5.20-5.28(m, 1H),
5.36-5.44(m, 1H), 6.90(br-s, 1H)
MS : 357.24(ES+)

30

Example 6-2

tert-Butyl (1S)-2-[(2S)-2-cyano-1-pyrrolidinyl]-
1-[[[(1-methylethylidene)amino]oxy]methyl]-2-oxoethyl
carbamate

5

To a solution of tert-butyl
(1S)-2-[(2S)-2-carbamoyl-1-pyrrolidinyl]-1-[[[(1-meth
ylethylidene)amino]oxy]methyl]-2-oxoethylcarbamate
(145mg) obtained by Example 6-1 in tetrahydrofuran (3ml)
10 was added triethylamine (0.17ml), then trifluoroacetic
anhydride (TFAA, 0.115ml) dropwise at 5°C. The reaction
mixture was stirred at 5°C for 30min.

Sat. NaHCO₃ was added and extracted with ethyl acetate.
The organic layer was dried over MgSO₄. After removal
15 of the solvent, the residue was dissolved in
isopropylether, washed with 1N HCl, sat. NaHCO₃, brine,
and dried over MgSO₄. After removal of the solvent in
vacuo, the residue was crystallized from n-hexane. The
crystals were collected by filtration, washed with
20 n-hexane to give the target compound as a white powder
(100mg, 72.6%).

¹H NMR(CDCl₃) : 1.43(s, 9H), 1.85(s, 3H), 1.88(s, 3H),
2.12-2.30(m, 4H), 3.62-3.84(m, 2H), 4.18-4.21(m, 2H),
25 4.67-4.81(m, 2H), 5.30-5.37(m, 1H)
MS : 339.19(ES+)

Example 6-3

(2S)-1-[(2S)-2-Amino-3-[[[(1-methylethylidene)amino]ox
30 y]propanoyl]-2-cyanopyrrolidine trifluoroacetate

tert-Butyl (1S)-2-[(2S)-2-cyano-1-pyrrolidinyl]-
1-[[[(1-methylethylidene)amino]oxy]methyl]-2-oxoethyl
carbamate (50mg) obtained by Example 6-2 was dissolved
5 in trifluoroacetic acid (1ml) at room temperature.

After 30min, the solvent was removed in vacuo, the
residue was crystallized from isopropylether, collected
by filtration, washed with isopropylether to give the
target compound as a white powder (37.3mg, 71.7%).

10

^1H NMR(DMSO- d_6) : 1.82(s, 3H), 1.83(s, 3H), 1.94-2.30(m,
4H), 3.53-3.72(m, 2H), 4.21-4.34(m, 2H), 4.40-4.46(m, 1H),
4.80-4.85(m, 1H), 8.34(br-s, 2H)

MS : 239.18(ES+), free 238.29

15

Example 7-1

tert-Butyl 4-[(benzyloxy)iminol]cyclohexylcarbamate

To a solution of tert-butyl 4-oxocyclohexylcarbamate
20 (300 mg) and triethylamine (0.235ml) in methanol (5ml)
at 0 °C was added O-benzylhydroxylamine hydrochloride
(225mg) at the same temperature. The resulting mixture
was warmed to room temperature and stirred for 2hrs.

The reaction was quenched with pouring H_2O . The
25 aqueous layer was extracted three times with ethyl acetate.
The combined organic layer was washed with brine, dried
over Na_2SO_4 , filtered and the solvent was removed in vacuo.
Recrystallization of the residue from hexane gave the
target compound (259mg, 57.9%).

30

¹H NMR (in CDCl₃) : 1.23-1.45(m, 12H), 1.94-2.16(m, 3H),
2.18-2.24(m, 1H), 2.37-2.45(m, 1H), 3.11-3.23(m, 1H),
3.76(br-s, 1H), 4.43(br-s, 1H), 5.06(s, 2H), 7.29-7.35(m,
5H)

5 MS : 319.20 (ES+)

Example 7-2

4-Aminocyclohexanone O-benzyloxime

10 To a solution of tert-butyl
4-[(benzyloxy)imino]cyclohexylcarbamate (32mg)
obtained by Example 7-1 in methanol (2ml) at room
temperature was added 4N HCl in dioxane (5ml).

After stirring for 15min, the reaction was quenched
15 with saturated aqueous solution of NaHCO₃. The aqueous
layer was extracted three times with CHCl₃. The combined
organic layer was washed with brine, dried over Na₂SO₄
and filtered. After removal of the solvent in vacuo, the
target compound was given as colorless oil (26mg, about
20 100%).

Example 7-3

(2S)-1-[[[4-[(Benzyloxy)imino]cyclohexyl]amino]acetyl
]-2-cyanopyrrolidine hydrochloride

25

To a solution of
(2S)-1-(bromoacetyl)-2-pyrrolidinecarbonitrile(32mg)
in tetrahydrofuran (2ml) at 0°C was added a solution of
4-aminocyclohexanone O-benzyloxime (64mg) obtained by
30 Example 7-2 in tetrahydrofuran (1ml) at that temperature

and stirred for 2hrs.

The reaction was quenched with pouring H₂O. The aqueous layer was extracted three times with ethyl acetate. The combined organic layer was washed with brine, dried
5 over Na₂SO₄, filtered and the solvent was removed in vacuo. The residue was purified with silica gel chromatography (CHCl₃:methanol=15:1). After removal of the solvent in vacuo, the residual oil was dissolved in methanol and 4N HCl-dioxane was added. After removal of the solvent, the
10 residue was triturated by diethylether to give the target compound as a white powder (26mg, 45.1%).

¹H NMR (in DMSO-d₆) : 1.39-1.65(m, 2H), 1.80-2.42(m, 9H),
3.11-3.41(m, 3H), 3.41-3.57(m, 1H), 3.59-3.73(m, 1H),
15 3.94-4.19(m, 2H), 5.34(dd, J=4.5, 6.6Hz, 1H), 5.01(s, 2H),
7.29-7.41(m, 5H), 9.27(br-s, 1H)
MS : 355.22 (ES+)

Example 8-1

20 tert-Butyl 4-[(2-pyridinylmethoxy)imino]cyclohexyl-
carbamate

To a solution of tert-butyl 4-oxocyclohexylcarbamate (100mg) in methanol (2ml) at room temperature was added
25 O-(2-pyridinylmethyl)hydroxylamine (58mg) at that temperature.

After stirring for 10 min, the solvent was removed in vacuo. The resulting precipitate was triturated with diethylether to give the target compound as white powder
30 (126mg, 84.1%).

¹H NMR (in CDCl₃) : 1.22-1.52(m, 11H), 1.96-2.28(m, 3H),
2.34-2.48(m, 1H), 3.15-3.30(m, 1H), 3.60-3.79(m, 1H),
4.36-4.55(m, 1H), 5.19(s, 2H), 7.15-7.22(m, 1H),
5 7.31-7.37(m, 1H), 7.64-7.72(m, 1H), 8.54-8.60(m, 1H)
MS : 320.18 (ES+)

Example 8-2

4-Aminocyclohexanone O-(2-pyridinylmethyl)oxime

10

Reaction was carried out in a manner similar to Example
7-2 using tert-butyl
4-[(2-pyridinylmethoxy)imino]cyclohexylcarbamate
obtained by Example 8-1 to give 80mg of target compound
15 (96.3%).

¹H NMR (in CDCl₃) : 1.21-1.55(m, 1H), 1.89-2.07(m, 2H),
2.07-2.23(m, 1H), 2.37-2.50(m, 1H), 2.91-3.06(m, 1H),
3.19-3.34(m, 1H), 3.71(s, 2H), 5.20(s, 2H), 7.12-7.24(m,
20 1H), 7.30-7.41(m, 1H), 7.62-7.73(m, 1H), 8.52-8.62(m, 1H)
MS : 220.17, 439.17 (ES+)

Example 8-3

(2S)-1-[[[4-[(2-Pyridinylmethoxy)imino]cyclohexyl]ami
25 no]acetyl]-2-cyanopyrrolidine hydrochloride

Reaction was carried out in a manner similar to Example
7-3 using 4-aminocyclohexanone
O-(2-pyridinylmethyl)oxime obtained by Example 8-2 to
30 give 17.2mg of target compound (22.7%).

¹H NMR (in DMSO-d₆) : 1.47-1.68(m, 2H), 1.87-2.40(m, 9H),
3.16(s, 2H), 3.20-3.38(m, 1H), 3.43-3.57(m, 1H),
3.61-3.73(m, 1H), 3.96-4.20(m, 2H), 4.81-4.90(m, 1H),
5 5.28(s, 2H), 7.69-7.82(m, 2H), 8.23-8.34(m, 1H),
8.72-8.79(m, 1H), 9.15-9.47(m, 2H)
MS 356.16 (ES+)

Example 9-1

10 tert-Butyl 4-[(3-pyridinylmethoxy)imino]cyclohexyl-
carbamate

Reaction was carried out in a manner similar to Example
8-1 using O-(3-pyridinylmethyl)hydroxylamine to give
15 180mg of target compound (60.1%).

¹H NMR (in CDCl₃) : 1.22-1.52(m, 11H), 1.91-2.16(m, 2H),
2.16-2.26(m, 1H), 2.32-2.46(m, 1H), 3.06-3.21(m, 1H),
3.56-3.78(m, 1H), 4.33-4.53(m, 1H), 5.06(s, 2H),
20 7.24-7.31(m, 1H), 7.63-7.70(m, 1H), 8.52-8.56(m, 1H),
8.58-8.61(m, 1H)
MS : 320.15 (ES+)

Example 9-2

25 4-Aminocyclohexanone O-(3-pyridinylmethyl)oxime

Reaction was carried out in a manner similar to Example
7-2 using tert-butyl
4-[(3-pyridinylmethoxy)imino]cyclohexylcarbamate
30 obtained by Example 9-1 to give 120mg of target compound

(97.1%).

MS : 220.17 (ES+) .

5 Example 9-3

(2S)-1-[[[4-[(3-Pyridinylmethoxy)imino]cyclohexyl]amino]acetyl]-2-cyanopyrrolidine hydrochloride

Reaction was carried out in a manner similar to Example
10 7-3 using 4-aminocyclohexanone
O-(3-pyridinylmethyl)oxime obtained by Example 9-2 to
give 49.1mg of target compound (37.3%).

¹H NMR (in DMSO-d₆) : 1.44-1.71(m, 2H), 1.86-2.42(m, 9H),
15 3.17-3.57(m, 2H), 3.57(s, 2H), 3.60-3.75(m, 1H),
3.94-4.21(m, 2H), 4.85(dd, J=4.2, 6.6Hz, 1H), 5.23(s, 2H),
8.00-8.09(m, 1H), 8.46-8.55(m, 1H), 8.85-8.92(m, 2H),
9.35(br-s, 1H), 9.46(br-s, 1H)
MS 356.19 (MS+)

20

Example 10-1

tert-Butyl 4-(methoxyimino)cyclohexylcarbamate
hydrochloride

25 Reaction was carried out in a manner similar to Example
7-1 using methoxyamine hydrochloride to give 224mg of
target compound (98.6%).

¹H NMR (in CDCl₃) : 1.20-1.53(m, 11H), 1.87-2.16(m, 4H),
30 2.16-2.27(m, 1H), 2.32-2.47(m, 1H), 2.99-3.17(m, 1H),

3.69(br-s, 1H), 3.82(s, 3H), 4.44(br-s, 1H)

MS : 243.17 (MS+)

Example 10-2

5 4-Aminocyclohexanone O-methyloxime

Reaction was carried out in a manner similar to Example
7-2 using tert-butyl
4-(methoxyimino)cyclohexylcarbamate hydrochloride
10 obtained by Example 10-1 to give 105mg of target compound
(79.8%).

¹H NMR (in CDCl₃) : 5.34(s, 2H), 7.26-7.30(m, 1H),
7.72-7.83(m, 6H), 8.55-8.56(m, 1H)

15 MS : 142.95 (ES+)

Example 10-3

(2S)-1-[[[4-(Methoxyimino)cyclohexyl]amino]acetyl]-2-
cyanopyrrolidine hydrochloride

20

Reaction was carried out in a manner similar to Example
7-3 using 4-aminocyclohexanone O-methyloxime obtained by
Example 10-2 to give 15mg of target compound (12.9%).

25 ¹H NMR (in DMSO-d₆) : 1.37-1.68(m, 2H), 1.76-2.41(m, 9H),
3.04-3.15(m, 1H), 3.19-3.77(m, 7H), 3.94-4.21(m, 2H),
4.81-4.98(m, 1H), 9.26(br-s, 1H)

MS : 279.13 (ES+)

30 Example 11-1

tert-Butyl 4-(tert-butoxyimino)cyclohexylcarbamate

Reaction was carried out in a manner similar to Example
7-1 using tert-butoxyamine hydrochloride to give 264mg
5 of target compound (99%).

MS : 285.25 (ES+)

Example 11-2

10 4-Aminocyclohexanone O-(tert-butyl)oxime

Reaction was carried out in a manner similar to Example
7-2 using tert-butyl
4-(tert-butoxyimino)cyclohexylcarbamate obtained by
15 Example 11-1 to give 120mg of target compound (70.1%).

MS : 185.13 (ES+)

Example 11-3

20 (2S)-1-[[[4-(tert-Butoxyimino)cyclohexyl]amino]acetyl
]-2-cyanopyrrolidine hydrochloride

Reaction was carried out in a manner similar to Example
7-3 using 4-aminocyclohexanone O-(tert-butyl)oxime
25 obtained by Example 11-2 to give 24.3mg of target compound
(21.1%).

¹H NMR (in DMSO-d₆) : 1.21(s, 9H), 1.35-1.61(m, 2H),
1.72-1.90(m, 1H), 1.92-2.44(m, 8H), 3.05-3.82(m, 6H),
30 3.96-4.22(m, 2H), 4.81-4.90(m, 1H), 9.20(br-s, 1H)

MS : 321.19 (ES+)

Example 12-1

tert-Butyl cis-4-[[[(5-cyano-2-pyridinyl)-
5 methylidene]amino]oxy]cyclohexylcarbamate

Reaction was carried out in a manner similar to Example 3-1 using 5-cyano-2-pyridinecarboxaldehyde to give 311mg of target compound (34.7%).

10

¹H NMR (in CDCl₃) : 1.34-2.10(m, 17H), 3.56(br-s, 1H),
4.34-4.57(m, 2H), 7.90-8.00(m, 2H), 8.18(s, 1H),
8.35-8.89(m, 1H)

15 Example 12-2

cis-6-[[[(4-Aminocyclohexyl)oxy]imino]methyl]nicotino
nitrile

Reaction was carried out in a manner similar to Example
20 3-2 using tert-butyl
cis-4-[[[(5-cyano-2-pyridinyl)methylidene]amino]oxy]c
yclohexylcarbamate obtained by Example 12-1 to give 265mg
of target compound (99.4%).

25 ¹H NMR (in DMSO-d₆) : 1.47-1.82(m, 6H), 1.99-2.14(m, 2H),
3.02-3.46(m, 2H), 4.43(br-s, 2H), 7.93-8.01(m, 1H),
8.26(s, 1H), 8.32-8.40(m, 6H), 9.05-9.10(m, 1H)
MS : 245.15 (ES+)

30 Example 12-3

cis-6-[[[4-[[2-[(2S)-2-Cyano-1-pyrrolidinyl]-2-oxoethyl]amino]cyclohexyl]oxy]imino]methyl]nicotinonitrile

Reaction was carried out in a manner similar to Example 7-3 using cis-6-[[[4-aminocyclohexyl]oxy]imino]methyl]nicotinonitrile obtained by Example 12-2 to give 68.1mg of target compound (38.9%).

¹H NMR (in CDCl₃) : 1.42-1.84(m, 8H), 2.00-2.37(m, 4H), 2.52-2.66(m, 1H), 3.43(s, 2H), 3.46-3.73(m, 2H), 4.37-4.48(m, 2H), 4.73-4.81(m, 1H), 7.26-7.30(m, 1H), 7.91-7.96(m, 2H), 8.18(s, 1H), 8.84-8.87(m, 1H)
MS : 381.19 (ES+)

15

Example 13-1

tert-Butyl trans-4-[(1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl)oxy]cyclohexylcarbamate

To a solution of tert-butyl cis-4-hydroxycyclohexylcarbamate (2g), hydroxyphthalimide (1.52g) and triphenylphosphine (2.68g) in tetrahydrofuran (50ml) at room temperature was added diisopropyl azodicarboxylate (DIAD, 2ml) and stirred at that temp for 2hrs.

The reaction was quenched by pouring H₂O. The aqueous layer was extracted with ethyl acetate. The combined organic layer was washed with brine, dried over Na₂SO₄, and filtered. After removal of the solvent in vacuo, the residue was triturated diethylether and a few drop of

hexane to give the target compound as a white powder (3.3g, 98.6%).

MS : 359.18 (ES-)

5

Example 13-2

tert-Butyl trans-4-(aminooxy)cyclohexylcarbamate

A solution of tert-butyl
10 trans-4-[(1,3-dioxo-1,3-dihydro-2H-isoindol-2-yl)oxy]
cyclohexylcarbamate (2g) obtained by Example 13-1 in
ethanol (200ml) was warmed to 80°C. After the solution
was turned clear, hydrazine monohydrate (9ml) was added
at that temperature. The resulting mixture was heated
15 at reflux for 1hr and then was cooled to room temperature.

The resulting precipitate was filtered off and washed
with ethyl acetate and the filtrate was concentrated. The
resulting mixture was diluted with 1N HCl and ethyl acetate,
and the water layer was extracted with ethyl acetate. The
20 aqueous layer was basified with solid NaHCO₃, saturated
with NaCl and was extracted three times with ethyl acetate.
The combined organic layer was washed with brine, dried
over Na₂SO₄, and filtered. After removal of the solvent
in vacuo, the target compound was given as a white powder
25 (850mg, 39.7%).

¹H NMR (in CDCl₃) : 1.03-1.37(m, 4H), 1.44(s, 9H),
1.37-1.94(m, 4H), 1.94-2.13(m, 1H), 3.33-3.53(m, 2H),
4.37(br-s, 1H), 5.29(br-s, 2H)

30 MS : 231.19 (ES+)

Example 13-3

trans-tert-Butyl 4-[[[(2-pyridinylmethylidene)amino]oxy]cyclohexylcarbamate

5

To a solution of tert-butyl trans-4-(aminooxy)cyclohexylcarbamate (250mg) obtained by Example 13-2 in methanol(2ml) at room temperature was added 2-pyridinecarboxaldehyde at that temperature.

10 After stirring for 10 min, the solvent was removed in vacuo. The resulting precipitate was triturated with diisopropylether to give the target compound as white powder (250mg, 72.1%).

15 ¹H NMR (in CDCl₃) : 1.14-1.34(m, 2H), 1.39-1.58(m, 11H), 2.00-2.25(m, 4H), 3.40-3.60(m, 1H), 4.12-4.27(m, 1H), 4.33-4.48(m, 1H), 7.21-7.29(m, 1H), 7.64-7.80(m, 2H), 8.15(s, 1H), 8.59-8.61(m, 1H)
MS : 320.18 (ES+)

20

Example 13-4

2-Pyridinecarbaldehyde O-(trans-4-aminocyclohexyl)-oxime

25 Reaction was carried out in a manner similar to Example 7-2 using trans-tert-butyl 4-[[[(2-pyridinylmethylidene)amino]oxy]cyclohexylcarbamate obtained by Example 13-3 to give 125mg of target compound (72.8%).

30

MS : 220.17 (ES+)

Example 13-5

trans-(2S)-1-[[[4-[[[(2-Pyridinylmethylidene)amino]oxy
5]cyclohexyl]amino]acetyl]-2-cyanopyrrolidine
dihydrochloride

Reaction was carried out in a manner similar to Example
7-3 using 2-pyridinecarbaldehyde
10 O-(trans-4-aminocyclohexyl)oxime obtained by Example
13-4 to give 67.5mg of target compound (55.2%).

Example 14-1

tert-Butyl trans-4-[[[(3-pyridinylmethylidene)amino]-
15 oxy]cyclohexyl]carbamate

Reaction was carried out in a manner similar to Example
13-3 using nicotinaldehyde to give 218mg of target
compound (78.6%).

20

¹H NMR (in CDCl₃) : 1.36-1.34(m, 2H), 1.38-1.57(m, 11H),
1.98-2.27(m, 4H), 3.41-3.59(m, 1H), 4.04-4.21(m, 1H),
4.32-4.48(m, 1H), 7.22-7.35(m, 1H), 7.86-7.98(m, 1H),
8.06(s, 1H), 8.54-8.63(m, 1H), 8.70-8.76(m, 1H)

25 MS : 318.30 (ES-)

Example 14-2

Nicotinaldehyde O-(trans-4-aminocyclohexyl)oxime

30 Reaction was carried out in a manner similar to Example

7-2 using tert-butyl
trans-4-[[[(3-pyridinylmethylidene)amino]oxy]cyclohexyl]carbamate obtained by Example 14-1 to give 155mg of target compound (about 100%).

5

Example 14-3

trans-(2S)-1-[[[4-[[[(3-Pyridinylmethylidene)amino]oxy]cyclohexyl]amino]acetyl]-2-cyanopyrrolidine

10 Reaction was carried out in a manner similar to Example
7-3 using nicotinaldehyde
O-(trans-4-aminocyclohexyl)oxime obtained by Example
14-2 to give 91.6mg of target compound (74.6%).

15 ¹H NMR (in CDCl₃) : 1.16-1.55(m, 4H), 1.91-2.05(m, 2H),
2.06-2.38(m, 6H), 2.42-2.61(m, 1H), 3.36-3.80(m, 5H),
4.09-4.26(m, 1H), 4.72-4.83(m, 1H), 7.23-7.34(m, 1H),
7.91-8.00(m, 1H), 8.07(s, 1H), 8.56-8.61(m, 1H),
8.70-8.74(m, 1H)

20 MS : 356.19 (ES+)

Example 15-1

tert-Butyl trans-4-[[[1-(3-pyridinyl)ethylidene]-
amino]oxy]cyclohexylcarbamate

25

Reaction was carried out in a manner similar to Example
13-3 using 3-acetylpyridine to give 166mg of target
compound (57.3%).

30 ¹H NMR (in CDCl₃) : 1.14-1.35(m, 2H), 1.39-1.59(m, 11H),

1.99-2.21(m, 4H), 2.23(s, 3H), 3.40-3.62(m, 1H),
4.06-4.22(m, 1H), 4.33-4.49(m, 1H), 7.23-7.32(m, 1H),
7.90-7.98(m, 1H), 8.54-8.59(m, 1H), 8.83-8.88(m, 1H)

MS : 334.23 (ES+)

5

Example 15-2

1-(3-Pyridinyl)ethanone O-(trans-4-aminocyclohexyl)-
oxime

Reaction was carried out in a manner similar to Example
7-2 using tert-butyl
trans-4-[[[[1-(3-pyridinyl)ethylidene]amino]oxy]cyclohexylcarbamate obtained by Example 15-1 to give 110mg of
target compound (94.7%).

15

Example 15-3

trans-(2S)-1-[[[4-[[[1-(3-Pyridinyl)ethylidene]amino]oxy]cyclohexyl]amino]acetyl]-2-cyanopyrrolidine dihydrochloride

20

Reaction was carried out in a manner similar to Example 7-3 using 1-(3-pyridinyl)ethanone O-(trans-4-aminocyclohexyl)oxime obtained by Example 15-2 to give 63mg of target compound (72%).

25

$$^1\text{H NMR (in DMSO-}d_6\text{)} : 1.34\text{--}2.42(\text{m}, 12\text{H}), 3.04\text{--}3.25(\text{m}, 1\text{H}),$$
$$3.33\text{--}3.96(\text{m}, 4\text{H}), 4.07\text{--}4.32(\text{m}, 2\text{H}), 4.90\text{--}4.99(\text{m}, 1\text{H}),$$
$$7.88\text{--}7.98(\text{m}, 1\text{H}), 8.54\text{--}8.66(\text{m}, 1\text{H}), 8.85\text{--}8.94(\text{m}, 1\text{H}),$$
$$9.08\text{--}9.14(\text{m}, 1\text{H}), 9.16\text{--}9.36(\text{m}, 1\text{H})$$

30 MS : 370.24 (ES+)

Example 16-1

tert-Butyl trans-4-[[[(5-cyano-2-pyridinyl)-
methylenediamino]oxy]cyclohexyl]carbamate

5

Reaction was carried out in a manner similar to Example 13-3 using 5-cyano-2-pyridinecarbaldehyde to give 190mg of target compound (62%).

10 ^1H NMR (in CDCl_3) : 5.34(s, 2H), 7.26-7.30(m, 1H),
7.72-7.83(m, 6H), 8.55-8.56(m, 1H)

Example 16-2

trans-6-[[[(4-Aminocyclohexyl)oxy]iminomethyl]nicotino
15 nitrile

To tert-butyl trans-4-[[[(5-cyano-2-pyridinyl)-
methylenediamino]oxy]cyclohexyl]carbamate (190mg)
obtained by Example 16-1 was added trifluoroacetic acid
20 (2ml) at room temperature and stirred for 2min.

The reaction was quenched with pouring a saturated
aqueous solution of NaHCO_3 . The aqueous layer was
extracted three times with CHCl_3 . The combined organic
layer was washed with brine, dried over Na_2SO_4 , and filtered.
25 The solvent was removed in vacuo to give the target compound
as white powder (135mg, about 100%).

^1H NMR (in CDCl_3) : 1.10-1.63(m, 13H), 1.99-2.24(m, 4H),
3.39-3.58(m, 1H), 4.12-4.30(m, 1H), 4.30-4.49(m, 1H),
30 7.87-7.97(m, 2H), 8.12-8.19(m, 1H), 8.80-8.90(m, 1H)

Example 16-3

trans-6-[[[4-[[2-[(2S)-2-Cyano-1-pyrrolidinyl]-2-oxoethyl]amino]cyclohexyl]oxy]iminomethyl]nicotinonitrile
5 dihydrochloride

Reaction was carried out in a manner similar to Example
7-3 using trans-6-[[[4-aminocyclohexyl]oxy]-
iminomethyl]nicotinonitrile obtained by Example 16-2 to
10 give 69mg of target compound (55.1%).

^1H NMR (in DMSO- d_6) : 0.948-2.29(m, 12H), 2.94-3.88(m,
5H), 3.92-4.26(m, 2H), 4.80-4.91(m, 1H), 7.92-7.99(m, 1H),
8.33-8.39(m, 1H), 9.01-9.26(m, 3H)
15 MS : 381.20 (ES+)

In order to illustrate the usefulness of the object
Compound (I), the pharmacological test is carried out as
shown in the following.

20

[A] Inhibition test of human plasma DPP-IV :

(i) Material and Method :

The effect of test compounds on DPP-IV activity in
human plasma was evaluated with a modified version of the
25 assay described by Hughes et al (Biochemistry, 38,
pp11597-11603(1999)).

Briefly, 20 μL of human plasma were mixed with 20
 μL of 80mM MgCl_2 in assay buffer (25mM HEPES, 140mM NaCl,
1% RIA-grade BSA, pH7.8), and were incubated in a room
30 temperature for 60min. Then the reaction was initiated

by the addition of both 20 μ L of test compounds and 20 μ L of 0.2mM substrate (H-glycine-proline-AMC; AMC is 7-amino-4-methylcoumarine), they were dissolved in the assay buffer.

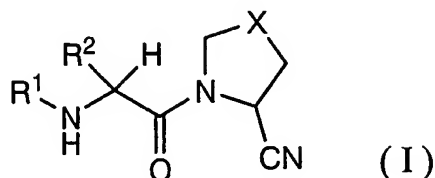
5 After 20min incubation in a room temperature (kept in the dark), fluorescence was measured (Excitation 380nm, Emission 460nm). A fluorescence-concentration curve of free AMC was obtained using AMC solution in the assay buffer with appropriate concentration. Plasma DPP-IV
10 activities, with or without the test compounds, were expressed as the amount of product per minute per mL. The potency of the test compounds as DPP-IV inhibitor was expressed as IC₅₀.

15 Industrial Applicability

 The Compound (I) or pharmaceutically acceptable salts thereof of the present invention have an inhibiting activity against DPP-IV. Therefore, the Compound (I) or pharmaceutically acceptable salts thereof are useful for
20 treating or preventing disease mediated by DPP-IV, more particularly useful for treating or preventing altered glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes mellitus (IDDM and NIDDM), diabetic neuropathy, nephropathy, and secondary diseases in
25 mammals or human caused by diabetes mellitus.

C L A I M S

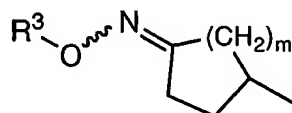
1. A compound of the formula (I) or pharmaceutically acceptable salt thereof.



[wherein

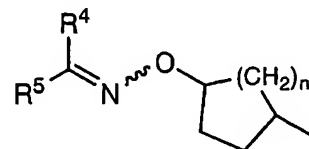
X is CH₂, O or S,

R¹ is hydrogen, lower alkyl, the moiety represented



by the formula:

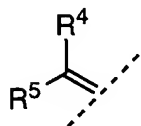
[wherein R³ is lower alkyl, aryl (which may have 1 to 3 substituent(s) selected from the group described later), heteroaryl (which may have 1 to 3 substituent(s) selected from the group described later), aryl-(lower alkyl) (which may have 1 to 3 substituent(s) selected from the group described later on the aryl group), or heteroaryl-(lower alkyl) (which may have 1 to 3 substituent(s) selected from the group described later on the heteroaryl group), m is integer of 1 to 3], or



the moiety represented by the formula:

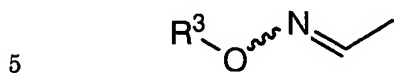
[wherein R⁴ is hydrogen or lower alkyl, R⁵ is lower alkyl, aryl (which may have 1 to 3 substituent(s) selected from the group described later), or heteroaryl (which may have 1 to 3 substituent(s) selected from the group

described later), or the partial structure:

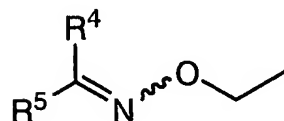


may form cycloalkyl, n is integer of 1 to 3],

R² is hydrogen, the moiety represented by the formula:



[wherein R³ represents the same definition defined above], or the moiety represented by the formula:

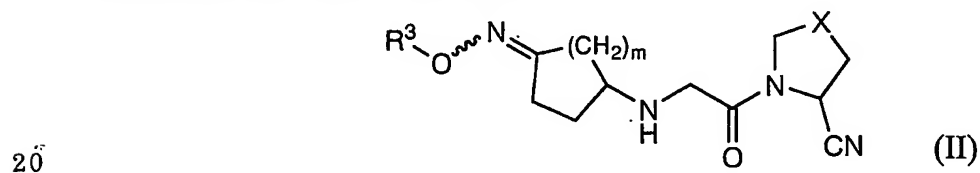


10 [wherein R⁴ and R⁵ represent the same definition defined above],

the substituent(s) is(are) selected from the group consisting of lower alkyl, lower alkoxy, halogen, cyano, nitro, amino and hydroxy,

15 provided that both of R¹ and R² are not hydrogen at the same time; and either R¹ and R² is hydrogen, or when R¹ is lower alkyl, R² is not hydrogen.]

2. A compound of the formula (II) or pharmaceutically acceptable salt thereof.



[wherein

X is CH₂, O or S,

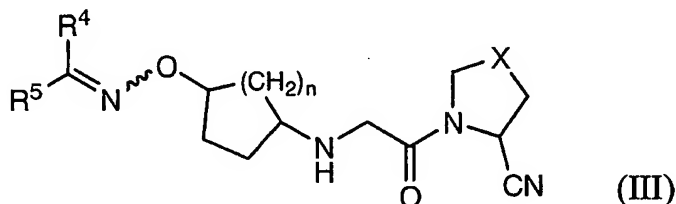
R³ is lower alkyl, aryl (which may have 1 to 3

substituent(s) selected from the group described later),
 heteroaryl (which may have 1 to 3 substituent(s) selected
 from the group described later), aryl-(lower alkyl) (which
 may have 1 to 3 substituent(s) selected from the group
 5 described later on the aryl group), or heteroaryl-(lower
 alkyl) (which may have 1 to 3 substituent(s) selected from
 the group described later on the heteroaryl group),

m is integer of 1 to 3

the substituent(s) is(are) selected from the group
 10 consisting of lower alkyl, halogen, cyano, nitro, amino
 and hydroxyl

3. A compound of the formula (III) or pharmaceutically
 acceptable salt thereof.



15

[wherein

X is CH_2 , O or S,

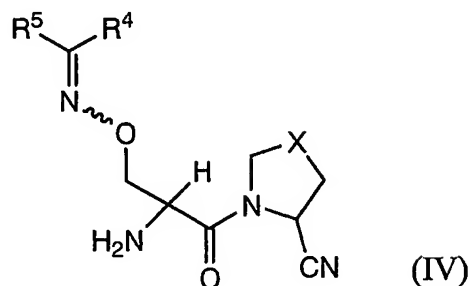
R^4 is hydrogen or lower alkyl,

R^5 is lower alkyl, aryl (which may have 1 to 3
 20 substituent(s) selected from the group described later),
 or heteroaryl (which may have 1 to 3 substituent(s)
 selected from the group described later),

n is integer of 1 to 3

the substituent(s) is(are) selected from the group
 25 consisting of lower alkyl, halogen, cyano, nitro, amino,
 and hydroxyl

4. A compound of the formula (IV) or pharmaceutically acceptable salt thereof.



[wherein

5 X is CH₂, O or S,

R⁴ is hydrogen or lower alkyl,

R⁵ is lower alkyl, aryl (which may have 1 to 3
substituent(s) selected from the group described later),
or heteroaryl (which may have 1 to 3 substituent(s)
10 selected from the group described later),

the substituent(s) is(are) selected from the group
consisting of lower alkyl, halogen, cyano, nitro, amino
and hydroxy]

15 5. A medicament comprising the compound according to any
one of Claim 1 to 4 as an active ingredient.

6. A pharmaceutical composition comprising the compound
according to any one of Claim 1 to 4 as an active ingredient,
20 in association with a pharmaceutically acceptable carrier
or excipient.

7. An inhibitor of DPP-IV consisting of the compound
according to any one of Claim 1 to 4.

25

8. A method for treatment and/or prevention of the medical conditions mediated by DPP-IV which comprises administering an effective amount of the compound according to any one of Claim 1 to 4 to human beings or
5 animals.

9. The method according to Claim 8, wherein the medical conditions mediated by DPP-IV is altered glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes
10 mellitus, diabetic neuropathy, nephropathy, or secondary diseases in mammals caused by diabetes mellitus.

10. The method according to Claim 8, wherein the medical conditions mediated by DPP-IV is NIDDM.
15

11. The compound according to any one of Claim 1 to 4 for use in the treatment and/or prevention of the medical conditions mediated by DPP-IV in human beings or animals.

20 12. The compound according to Claim 11, wherein the medical conditions mediated by DPP-IV is altered glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes mellitus, diabetic neuropathy, nephropathy, or secondary diseases in mammals caused by
25 diabetes mellitus.

13. The compound according to Claim 11, wherein the medical conditions mediated by DPP-IV is NIDDM.

30 14. Use of the compound according to any one of Claim

1 to 4 for the manufacture of a medicament for treatment and/or prevention of the medical conditions mediated by DPP-IV in human beings or animals.

5 15. The use according to Claim 14, wherein the medical conditions mediated by DPP-IV is altered glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes mellitus, diabetic neuropathy, nephropathy, or secondary diseases in mammals caused by diabetes mellitus.

10

16. The use according to Claim 14, wherein the medical conditions mediated by DPP-IV is NIDDM.

15 17. A commercial package comprising the pharmaceutical composition containing the compound identified in any one of Claim 1 to 4 and a written matter associated therewith, wherein the written matter states that the compound can or should be used for preventing and/or treating the medical conditions mediated by DPP-IV.

20

18. The commercial package according to Claim 17, wherein the medical conditions mediated by DPP-IV is altered glucose tolerance, glucosuria, hyperlipidemia, metabolic acidosis, diabetes mellitus, diabetic neuropathy, 25 nephropathy, or secondary diseases in mammals caused by diabetes mellitus.

19. The commercial package according to Claim 17, wherein the medical conditions mediated by DPP-IV is NIDDM.

30